

EAST Search History

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	26	(546/269.4).CCLS.	US-PGPUB	OR	OFF	2007/09/20 11:48

10/564,804

=> file casreact

FILE 'CASREACT' ENTERED AT 10:57:25 ON 20 SEP 2007

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FILE CONTENT:1840 - 15 Sep 2007 VOL 147 ISS 13

New CAS Information Use Policies, enter HELP USAGETERMS for details.

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*
*      CASREACT now has more than 12 million reactions      *
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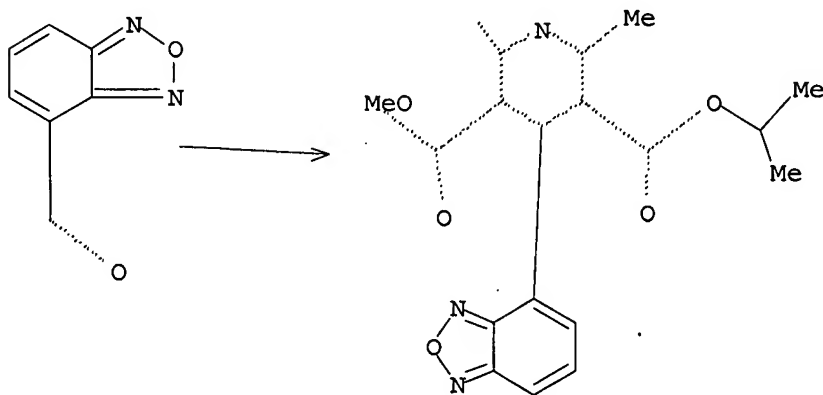
Some CASREACT records are derived from the ZIC/VINITI database (1974-1999) provided by InfoChem, INPI data prior to 1986, and Biotransformations database compiled under the direction of Professor Dr. Klaus Kieslich.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d que

L1

STR



Structure attributes must be viewed using STN Express query preparation.

L3

1 SEA FILE=CASREACT SSS FUL L1 (1 REACTIONS)

=> d l3 ibib abs fcrd

L3 ANSWER 1 OF 1 CASREACT COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER:

142:155958 CASREACT

TITLE:

Process for the manufacture of isradipine from the condensation reaction of 2,1,3-benzoxadiazole-4-carboxaldehyde and methyl acetoacetate and cyclocondensation of the intermediate with isopropyl β -aminocrotonate

INVENTOR(S):

Thakashinamoorthy, Chandiran; Senthil, Kumar Minor; Palanivel, Kaliyaperumal; Mullaiyur, Radhakrishnan Selvaraju

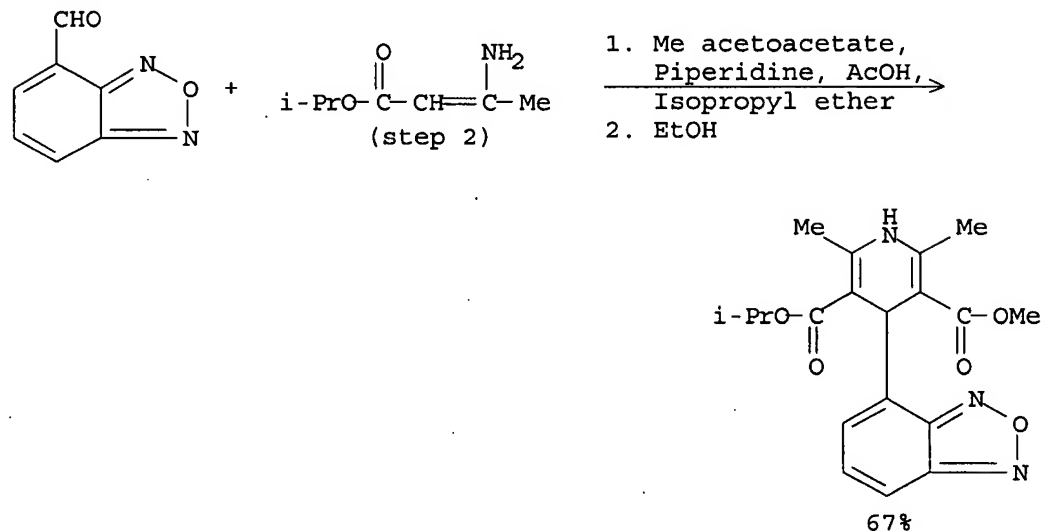
10/564,804

PATENT ASSIGNEE(S): Shasun Chemicals and Drugs Limited, India
 SOURCE: PCT Int. Appl., 14 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005005437	A1	20050120	WO 2004-IN208	20040715
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW. RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
IN 2003CH00571	A	20070504	IN 2003-CH571	20030715
US 2006167063	A1	20060727	US 2006-564804	20060113
PRIORITY APPLN. INFO.:			IN 2003-CH571	20030715
			WO 2004-IN208	20040715

AB Isradipine, 4-(4-benzofurazanyl)-1,4-dihydro-2,6-dimethyl-3,5-pyridinedicarboxylic acid Me 1-methylethyl ester (I), is prepared in high yield and selectivity by reacting 2,1,3-benzoxadiazole-4-carboxaldehyde with Me acetoacetate in the presence of acetic acid and piperidine in diisopropyl ether to obtain 2-acetyl-3-benzofurazan-4-ylacrylic acid Me ester which is purified and then reacted with iso-Pr β -aminocrotonate in ethanol at 25-35° to give I.

RX(3) OF 3 - 2 STEPS



CON: STEP(1.1) room temperature; 70 deg C
 STEP(2.1) room temperature; 5 hours, 25 - 28 deg C

REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

10/564,804

=> => file caplus

FILE 'CAPLUS' ENTERED AT 10:59:08 ON 20 SEP 2007

USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.

PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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Copyright of the articles to which records in this database refer is held by the publishers listed in the PUBLISHER (PB) field (available for records published or updated in Chemical Abstracts after December 26, 1996), unless otherwise indicated in the original publications. The CA Lexicon is the copyrighted intellectual property of the American Chemical Society and is provided to assist you in searching databases on STN. Any dissemination, distribution, copying, or storing of this information, without the prior written consent of CAS, is strictly prohibited.

FILE COVERS 1907 - 20 Sep 2007 VOL 147 ISS 13

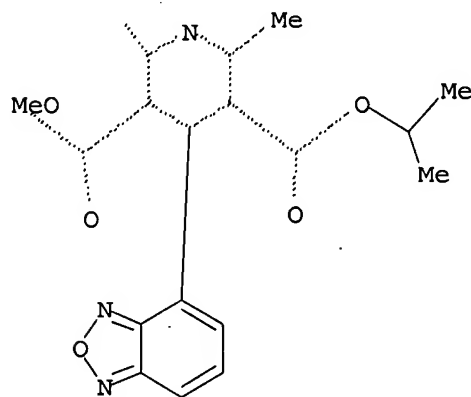
FILE LAST UPDATED: 19 Sep 2007 (20070919/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

<http://www.cas.org/infopolicy.html>

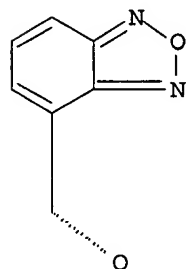
=> d que

L4 STR



Structure attributes must be viewed using STN Express query preparation.

L5 STR



Structure attributes must be viewed using STN Express query preparation.

L6 31 SEA FILE=REGISTRY SSS FUL L4

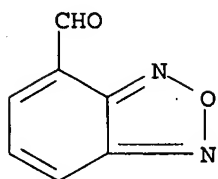
10/564,804

L7 42 SEA FILE=REGISTRY SSS FUL L5
L9 8 SEA FILE=CAPLUS L6 AND L7

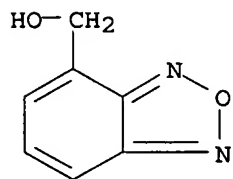
=> d 19 1-8 ibib abs hitstr

L9 ANSWER 1 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 2005:238972 CAPLUS
DOCUMENT NUMBER: 142:316844
TITLE: Process for the manufacture of 2,1,3-benzoxadiazole-4-carboxaldehyde
INVENTOR(S): Chandiran, Thakashina Moorthy; Subramaniam, S. S.; Swaminathan, Venkatraman
PATENT ASSIGNEE(S): Shasun Chemicals and Drugs Limited, India
SOURCE: PCT Int. Appl., 11 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

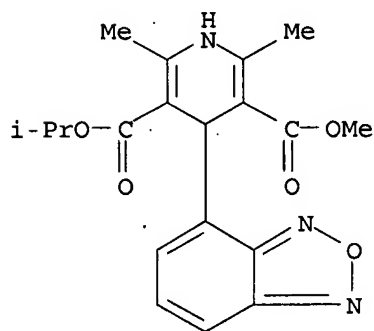
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005023787	A1	20050317	WO 2003-IN305	20030910
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2003269482	A1	20050329	AU 2003-269482	20030910
IN 2005CN02579	A	20070720	IN 2005-CN2579	20051010
IN 2005CN03165	A	20070727	IN 2005-CN3165	20051128
PRIORITY APPLN. INFO.:			WO 2003-IN305	A 20030910
OTHER SOURCE(S): CASREACT 142:316844				
AB A process is disclosed for the manufacture of 2,1,3-benzoxadiazole-4-carboxaldehyde, an intermediate for the preparation of 4-(4-Benzofurazanyl)-1,4-dihydro-2,6-dimethyl-3,5-pyridinedicarboxylic acid Me 1-methylethyl ester (Isradipine). For instance, 2,1,3-benzoxadiazole-4-yl-methanol is oxidized to 2,1,3-benzoxadiazole-4-carboxaldehyde using pyridinium chlorochromate (PCC) in CH ₂ Cl ₂ (0° (during addition of carbinol) -> 25-30°, 2 h) in 51% yield with 98.2% purity by HPLC (after silica gel filtration). The current process is more amenable to large scale preparation than prior art methods.				
IT 32863-32-4P, 2,1,3-Benzoxadiazole-4-carboxaldehyde RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation) (process for manufacture of 2,1,3-benzoxadiazole-4-carboxaldehyde)				
RN 32863-32-4 CAPLUS				
CN 2,1,3-Benzoxadiazole-4-carboxaldehyde (CA INDEX NAME)				



IT 175609-19-5, (2,1,3-Benzoxadiazole-4-yl)methanol
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (process for manufacture of 2,1,3-benzoxadiazole-4-carboxaldehyde)
 RN 175609-19-5 CAPLUS
 CN 2,1,3-Benzoxadiazole-4-methanol (CA INDEX NAME)



IT 75695-93-1, Isradipine
 RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (process for manufacture of 2,1,3-benzoxadiazole-4-carboxaldehyde)
 RN 75695-93-1 CAPLUS
 CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-2,6-dimethyl-, 3-methyl 5-(1-methylethyl) ester (CA INDEX NAME)



REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 2 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2005:58213 CAPLUS

DOCUMENT NUMBER: 142:155958

TITLE: Process for the manufacture of isradipine from the condensation reaction of 2,1,3-benzoxadiazole-4-carboxaldehyde and methyl acetoacetate and cyclocondensation of the intermediate with isopropyl β -aminocrotonate

INVENTOR(S): Thakashinamoorthy, Chandiran; Senthil, Kumar Minor; Palanivel, Kaliyaperumal; Mullaiyur, Radhakrishnan Selvaraju

PATENT ASSIGNEE(S): Shasun Chemicals and Drugs Limited, India

SOURCE: PCT Int. Appl., 14 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005005437	A1	20050120	WO 2004-IN208	20040715
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
IN 2003CH00571	A	20070504	IN 2003-CH571	20030715
US 2006167063	A1	20060727	US 2006-564804	20060113
PRIORITY APPLN. INFO.:			IN 2003-CH571	A 20030715
			WO 2004-IN208	W 20040715

OTHER SOURCE(S): CASREACT 142:155958

AB Isradipine, 4-(4-benzofurazanyl)-1,4-dihydro-2,6-dimethyl-3,5-pyridinedicarboxylic acid Me 1-methylethyl ester (I), is prepared in high yield and selectivity by reacting 2,1,3-benzoxadiazole-4-carboxaldehyde with Me acetoacetate in the presence of acetic acid and piperidine in diisopropyl ether to obtain 2-acetyl-3-benzofurazan-4-ylacrylic acid Me ester which is purified and then reacted with iso-Pr β -aminocrotonate in ethanol at 25-35° to give I.

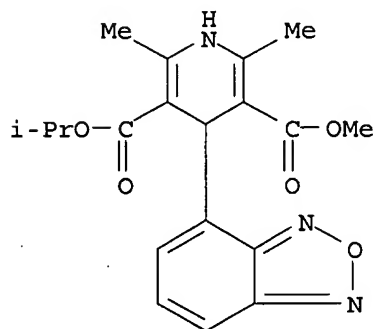
IT 75695-93-1P, Isradipine

RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)

(process for the manufacture of isradipine from the condensation reaction of 2,1,3-benzoxadiazole-4-carboxaldehyde and Me acetoacetate and cyclocondensation of the intermediate with iso-Pr β -aminocrotonate)

RN 75695-93-1 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-2,6-dimethyl-, 3-methyl 5-(1-methylethyl) ester (CA INDEX NAME)



IT 32863-32-4, 2,1,3-Benzoxadiazole-4-carboxaldehyde

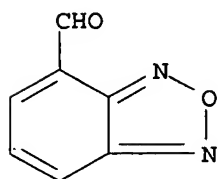
RL: RCT (Reactant); RACT (Reactant or reagent)

(process for the manufacture of isradipine from the condensation reaction of 2,1,3-benzoxadiazole-4-carboxaldehyde and Me acetoacetate and cyclocondensation of the intermediate with iso-Pr β -aminocrotonate)

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RN 32863-32-4 CAPLUS

CN 2,1,3-Benzoxadiazole-4-carboxaldehyde (CA INDEX NAME)



REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 3 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1994:217302 CAPLUS

DOCUMENT NUMBER: 120:217302

TITLE: Preparation of 1-(2-nitrobenzyl)-1,4-dihydropyridinecarboxylates as light-activated prodrugs for calcium modulators

INVENTOR(S): Goldmann, Siegfried; Bechem, Martin

PATENT ASSIGNEE(S): Bayer A.-G., Germany

SOURCE: Ger. Offen., 19 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent

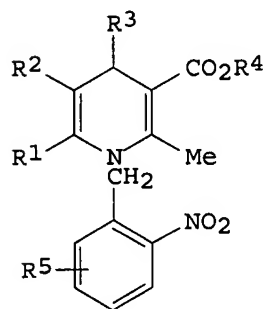
LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 4222770	A1	19940113	DE 1992-4222770	19920710
WO 9401405	A1	19940120	WO 1993-EP1720	19930703
W: JP, US				
RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
EP 650477	A1	19950503	EP 1993-915785	19930703
R: DE, FR, GB				
US 5606066	A	19970225	US 1995-356410	19950501
PRIORITY APPLN. INFO.:			DE 1992-4222770	A 19920710
			WO 1993-EP1720	W 19930703

OTHER SOURCE(S): MARPAT 120:217302
GI



I

AB Title compds. [I; R1 = H, cyano, CHO, alkyl, etc.; R2 = cyano, NO2, alkoxy carbonyl, etc.; R1R2 = CH2O2C; R3 = aryl, heterocyclyl, etc.; R4 = (substituted)alk(en)yl; R5 = H, halo, OH, CO2H, etc.] were prepared Thus, 3-(O2N)C6H4CHO was cyclocondensed with MeCOCH2CO2Me and 2-(O2N)C6H4CH2NH2

10/564,804

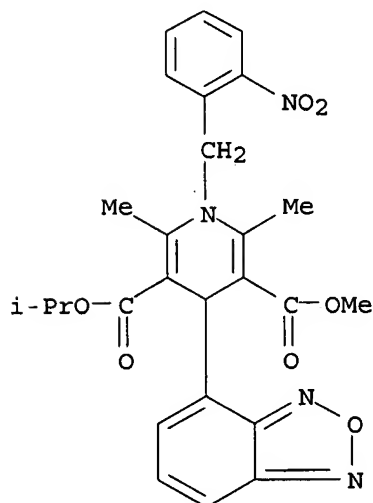
to give I [R1 = R4 = Me, R2 = CO2Me, R3 = 3-(O2N)C6H4, R5 = H].
Contractile force tracings of perfused guinea pig papillary muscle under
unirrad. and irradiation treatment were given.

IT 154026-72-9P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of, as photoactivated calcium modulator prodrug)

RN 154026-72-9 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-
2,6-dimethyl-1-[(2-nitrophenyl)methyl]-, methyl 1-methylethyl ester (9CI)
(CA INDEX NAME)



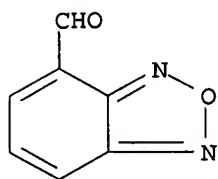
IT 32863-32-4, 2,1,3-Benzoxadiazole-4-carboxaldehyde

RL: RCT (Reactant); RACT (Reactant or reagent)

(reaction of, in preparation of photoactivated calcium modulator prodrug)

RN 32863-32-4 CAPLUS

CN 2,1,3-Benzoxadiazole-4-carboxaldehyde (CA INDEX NAME)



L9 ANSWER 4 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1991:185276 CAPLUS

DOCUMENT NUMBER: 114:185276

TITLE: A process for preparation of enantiomerically pure
polysubstituted 1,4-dihydropyridines

INVENTOR(S): Gandolfi, Carmelo A.; Frigerio, Marco; Riva, Carlo;
Zaliani, Andrea; Long, Giorgio; Di Domenico, Roberto

PATENT ASSIGNEE(S): Boehringer Biochemia Robin S.p.A., Italy

SOURCE: Eur. Pat. Appl., 32 pp.

CODEN: EPXXDW

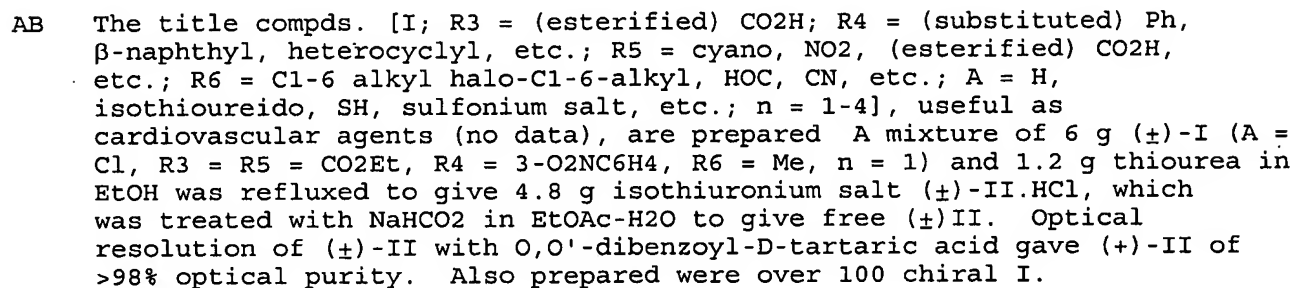
DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

OTHER SOURCE(S) : MARPAT 114:185276
GI



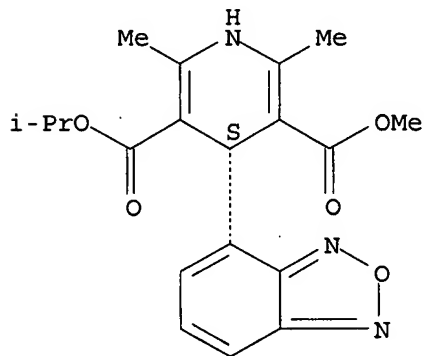
IT 84260-64-0P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of, as cardiovascular agent)

10/564,804

RN 84260-64-0 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-2,6-dimethyl-, methyl 1-methylethyl ester, (4S)- (CA INDEX NAME)

Absolute stereochemistry.



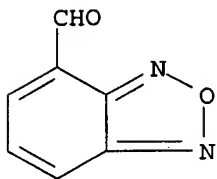
IT 32863-32-4, 4-Benzofurazancarboxaldehyde

RL: RCT (Reactant); RACT (Reactant or reagent)

(reaction of, with in preparation of cardiovascular agents)

RN 32863-32-4 CAPLUS

CN 2,1,3-Benzoxadiazole-4-carboxaldehyde (CA INDEX NAME)



L9 ANSWER 5 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1988:94566 CAPLUS

DOCUMENT NUMBER: 108:94566

TITLE: Preparation of benzoxa- and benzothiadiazoledicarboxaldehydes and their conversion to dihydropyridines

INVENTOR(S): Heitzmann, Markus

PATENT ASSIGNEE(S): Sandoz A.-G., Switz.

SOURCE: Patentschrift (Switz.), 4 pp.

CODEN: SWXXAS

DOCUMENT TYPE: Patent

LANGUAGE: German

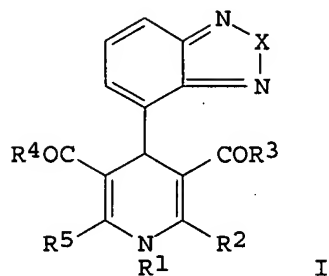
FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CH 661270	A5	19870715	CH 1982-6651	19821115
CH 661728	A5	19870814	CH 1982-1016	19821115
HU 44768	A2	19880428	HU 1986-1793	19860429
HU 196384	B	19881128		
DD 259400	A5	19880824	DD 1987-301996	19870421
PRIORITY APPLN. INFO.:			GB 1981-34708	A 19811118
			CH 1982-6651	A 19821115

OTHER SOURCE(S): CASREACT 108:94566

GI



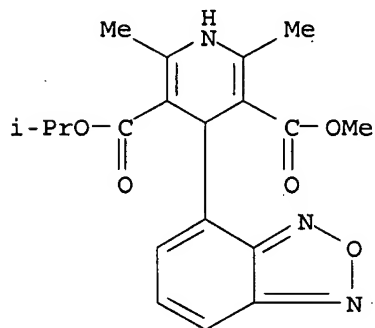
AB The title compds. [I; R1 = H, alkyl, alkenyl, alkynyl, cycloalkyl, (substituted) phenylalkyl; R2, R5 = H, alkyl; R3, R4 = alkyl, alkenyl, alkynyl, cycloalkyl, alkoxy, hydroxyalkoxy, alkenyloxy, etc; X = O, S] were prepared as drugs (no data). 2,1,3-Benzoxadiazole in THF was added to a solution of Li diisopropylamide in THF-cyclohexane at -75° and the resulting salt was added to DMF in THF. The initial adduct was hydrolyzed with aqueous HOAc and II was isolated via its bisulfite adduct. II was refluxed with Et acetoacetate and NH4OAc in EtOH to give I (R1 = H, R2 = R5 = Me, R3 = R4 = OEt).

IT 75695-93-1P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of, as drug)

RN 75695-93-1 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-2,6-dimethyl-, 3-methyl 5-(1-methylethyl) ester (CA INDEX NAME)

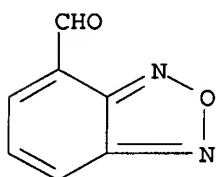


IT 32863-32-4P, 2,1,3-Benzoxadiazole-4-carboxaldehyde

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of, as drug intermediate)

RN 32863-32-4 CAPLUS

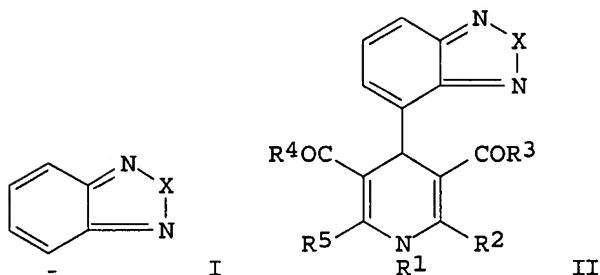
CN 2,1,3-Benzoxadiazole-4-carboxaldehyde (CA INDEX NAME)



L9 ANSWER 6 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1987:636717 CAPLUS
 DOCUMENT NUMBER: 107:236717
 TITLE: Preparation of metallated 2,1,3-benzoxa-or
 -thiadiazole and their use in synthesis of
 dihydropyridine derivatives
 INVENTOR(S): Heitzmann, Markus
 PATENT ASSIGNEE(S): Sandoz A.-G., Switz.
 SOURCE: Patentschrift (Switz.), 3 pp.
 CODEN: SWXXAS
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

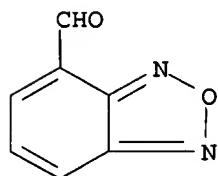
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CH 661728	A5	19870814	CH 1982-1016	19821115
CH 661270	A5	19870715	CH 1982-6651	19821115
PRIORITY APPLN. INFO.:			GB 1981-34708	A 19811118
			CH 1982-6651	A 19821115

GI



AB The title anions (I; X = O,S) were prepared by metalation of the parent heterocycles for use in synthesis of pharmacol. active dihydropyridines II [R1 = H, alkyl, alkenyl, alkynyl, cycloalkyl, (substituted) phenylalkyl; R2,R5 = H, alkyl; R3,R4 = alkyl, alkenyl, alkynyl, alkoxy, alkynyloxy, etc.]. 2,1,3-Benzoxadiazole in THF was added to a solution of LiN(CHMe2)2 in THF/cyclohexane at -78° and stirred for 35 min to afford the Li salt of I(X = O), which was added to DMF in THF followed by HOAc/H2O workup to give 2,1,3-benzoxadiazole-4-carboxaldehyde. The latter was cyclocondensed with MeCOCH2CO2Et and NH4OAc in EtOH to give II (R1 = H, R2 = R5 = Me, R3 = R4 = Et, X = O).

IT 32863-32-4
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (cyclocondensation of, with acetoacetate and ammonium acetate)
 RN 32863-32-4 CAPLUS
 CN 2,1,3-Benzoxadiazole-4-carboxaldehyde (CA INDEX NAME)



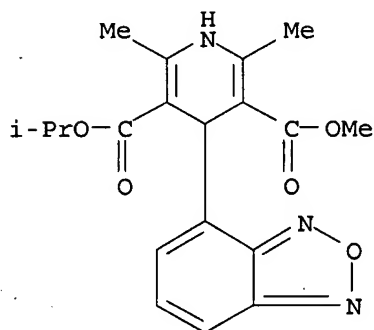
10/564,804

IT 75695-93-1P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of, as drug)

RN 75695-93-1 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-2,6-dimethyl-, 3-methyl 5-(1-methylethyl) ester (CA INDEX NAME)



L9 ANSWER 7 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1984:407162 CAPLUS

DOCUMENT NUMBER: 101:7162

TITLE: 1,4-Dihydropyridine derivatives in optically active or in racemate form and their pharmaceutical compositions

INVENTOR(S): Vogel, Arnold

PATENT ASSIGNEE(S): Sandoz-Patent-G.m.b.H., Fed. Rep. Ger.

SOURCE: Ger. Offen., 29 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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DE 3320616	A1	19831215	DE 1983-3320616	19830608
FR 2528431	A1	19831216	FR 1983-9551	19830607
FR 2528431	B1	19860110		
FI 8302072	A	19831216	FI 1983-2072	19830608
BE 897000	A1	19831209	BE 1983-10806	19830609
NL 8302067	A	19840102	NL 1983-2067	19830610
WO 8400033	A1	19840105	WO 1983-CH73	19830610
W: CH				
CH 660593	A5	19870515	CH 1984-688	19830610
DK 8302711	A	19831216	DK 1983-2711	19830613
GB 2122192	A	19840111	GB 1983-16054	19830613
GB 2122192	B	19851218		
CA 1208639	A1	19860729	CA 1983-430300	19830613
IL 68975	A	19870130	IL 1983-68975	19830613
SE 8303385	A	19831216	SE 1983-3385	19830614
AU 8315753	A	19831222	AU 1983-15753	19830614
JP 59005183	A	19840112	JP 1983-106597	19830614
HU 33802	A2	19841228	HU 1983-2113	19830614
HU 191853	B	19870428		
ES 523243	A1	19850401	ES 1983-523243	19830614
AT 8302183	A	19870315	AT 1983-2183	19830614
AT 384219	B	19871012		

10/564,804

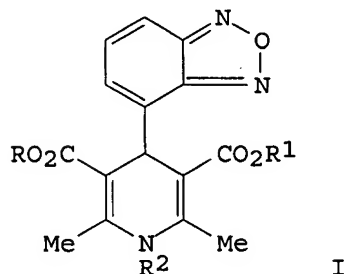
ZA 8304409
ES 538157
PRIORITY APPLN. INFO.:

A 19850227
A1 19860201

ZA 1983-4409
ES 1984-538157
CH 1982-3692
CH 1982-3693
WO 1983-CH73

19830615
19841130
A 19820615
A 19820615
A 19830610

OTHER SOURCE(S): MARPAT 101:7162
GI



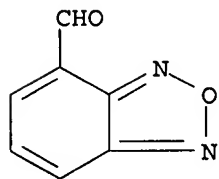
AB Calcium channel blocking (no data) benzoxadiazolylpyridinedicarboxylates I [1 of R, R1 = Me, the other = Me2CH, Bu, Me2CHCH2, Me2CHOCH2CH2, cyclopentyl; R2 = (un)substituted C1-6 alkyl] were prepared Thus, I (R = Me, R1 = Me2CH, R2 = H) was treated with (MeO)2SO2 in Me2SO to give (±)-I (R = Me, R1 = Me2CH, R2 = Me).

IT 32863-32-4

RL: RCT (Reactant); RACT (Reactant or reagent)
(condensation of, with acetoacetate derivative)

RN 32863-32-4 CAPLUS

CN 2,1,3-Benzoxadiazole-4-carboxaldehyde (CA INDEX NAME)

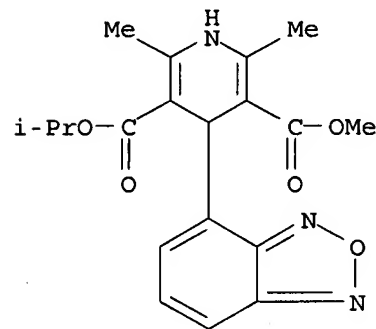


IT 75695-93-1

RL: RCT (Reactant); RACT (Reactant or reagent)
(methylation of)

RN 75695-93-1 CAPLUS

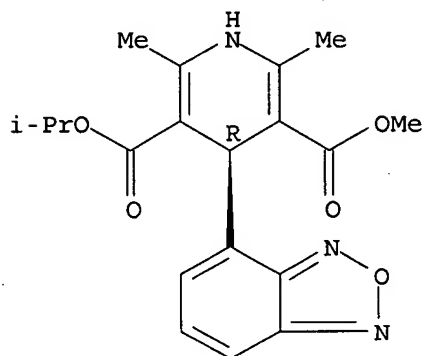
CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-2,6-dimethyl-, 3-methyl 5-(1-methylethyl) ester (CA INDEX NAME)



10/564,804

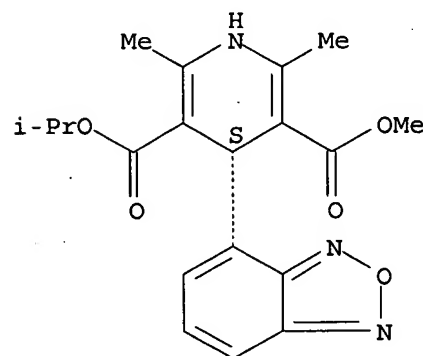
IT 84260-63-9P 84260-64-0P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
(Reactant or reagent)
(preparation and methylation of)
RN 84260-63-9 CAPLUS
CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-
2,6-dimethyl-, methyl 1-methylethyl ester, (4R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



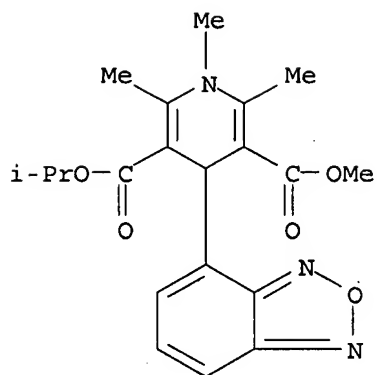
RN 84260-64-0 CAPLUS
CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-
2,6-dimethyl-, methyl 1-methylethyl ester, (4S)- (CA INDEX NAME)

Absolute stereochemistry.



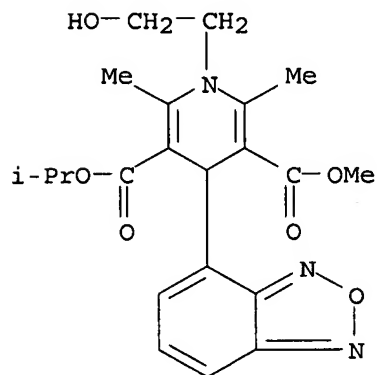
IT 88977-21-3P 88977-39-3P 88977-40-6P
88977-41-7P 88977-42-8P 88977-44-0P
88977-45-1P 88977-46-2P 88987-07-9P
88987-08-0P 88987-09-1P 89016-38-6P
89016-39-7P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
RN 88977-21-3 CAPLUS
CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-
1,2,6-trimethyl-, methyl 1-methylethyl ester (9CI) (CA INDEX NAME)

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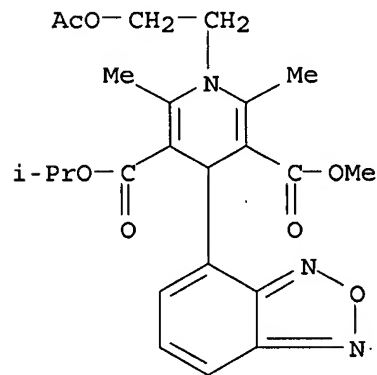
RN 88977-39-3 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-1-(2-hydroxyethyl)-2,6-dimethyl-, methyl 1-methylethyl ester (9CI) (CA INDEX NAME)



RN 88977-40-6 CAPLUS

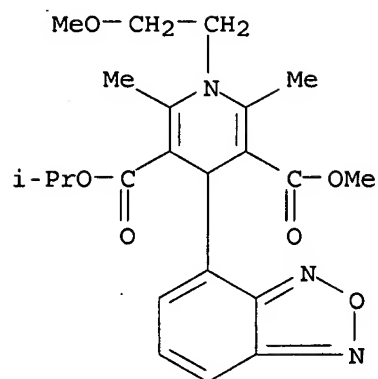
CN 3,5-Pyridinedicarboxylic acid, 1-[2-(acetyloxy)ethyl]-4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-2,6-dimethyl-, methyl 1-methylethyl ester (9CI) (CA INDEX NAME)



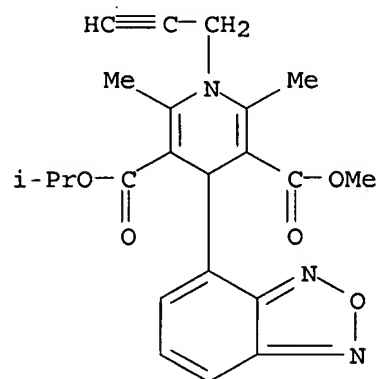
RN 88977-41-7 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-1-(2-methoxyethyl)-2,6-dimethyl-, methyl 1-methylethyl ester (9CI) (CA INDEX NAME)

10/564,804



RN 88977-42-8 CAPLUS
CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-2,6-dimethyl-1-(2-propynyl)-, methyl 1-methylethyl ester (9CI) (CA INDEX NAME)

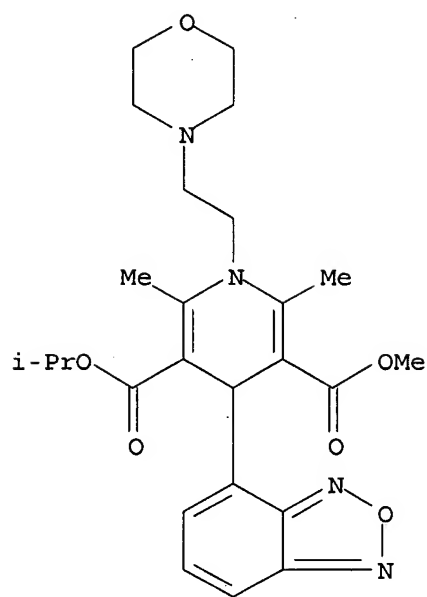


RN 88977-44-0 CAPLUS
CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-2,6-dimethyl-1-[2-(4-morpholinyl)ethyl]-, methyl 1-methylethyl ester, (2E)-2-butenedioate (1:1) (9CI) (CA INDEX NAME)

CM 1

CRN 88977-43-9
CMF C25 H32 N4 O6

10/564,804

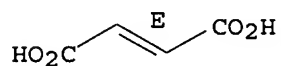


CM 2

CRN 110-17-8

CMF C4 H4 O4

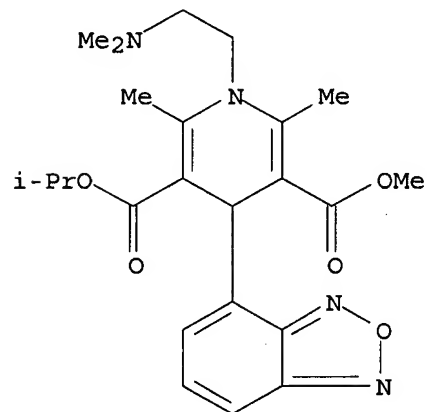
Double bond geometry as shown.



RN 88977-45-1 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1-[2-(dimethylamino)ethyl]-1,4-dihydro-2,6-dimethyl-, methyl 1-methylethyl ester, (-)-(9CI) (CA INDEX NAME)

Rotation (-).



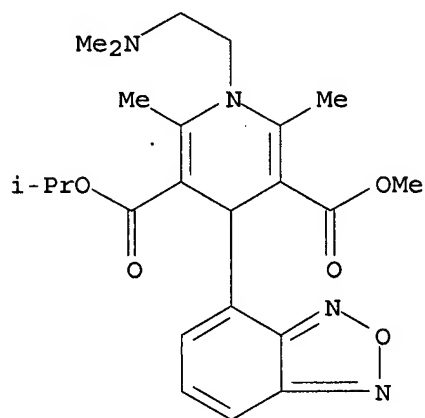
RN 88977-46-2 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1-[2-(dimethylamino)ethyl]-1,4-dihydro-2,6-dimethyl-, methyl 1-methylethyl

10/564,804

ester, hydrobromide, (-) - (9CI) (CA INDEX NAME)

Rotation (-).



●x HBr

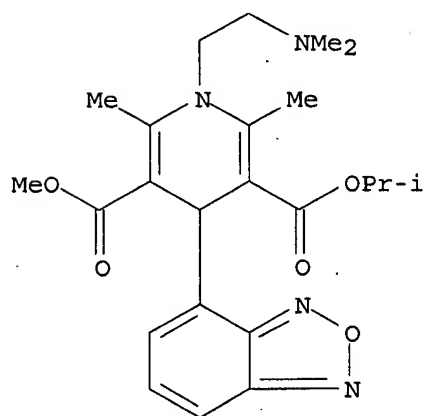
RN 88987-07-9 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1-[2-(dimethylamino)ethyl]-1,4-dihydro-2,6-dimethyl-, methyl 1-methylethyl ester, (2E)-2-butenedioate (1:1) (9CI) (CA INDEX NAME)

CM 1

CRN 88987-06-8

CMF C23 H30 N4 O5



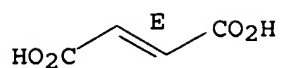
CM 2

CRN 110-17-8

CMF C4 H4 O4

Double bond geometry as shown.

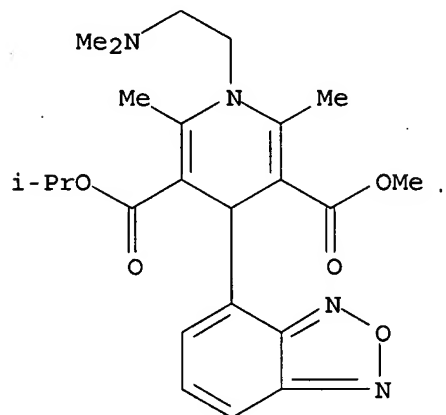
10/564,804



RN 88987-08-0 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1-[2-(dimethylamino)ethyl]-1,4-dihydro-2,6-dimethyl-, methyl 1-methylethyl ester, (+)-(9CI) (CA INDEX NAME)

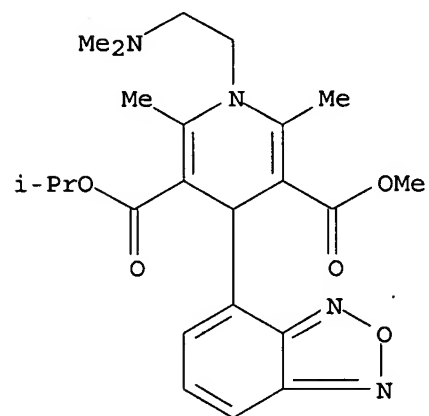
Rotation (+).



RN 88987-09-1 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1-[2-(dimethylamino)ethyl]-1,4-dihydro-2,6-dimethyl-, methyl 1-methylethyl ester, hydrobromide, (+)-(9CI) (CA INDEX NAME)

Rotation (+).



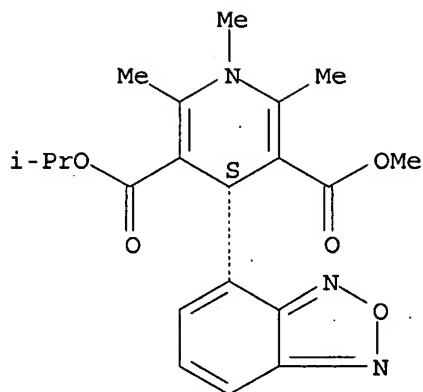
●x HBr

RN 89016-38-6 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-1,2,6-trimethyl-, methyl 1-methylethyl ester, (S)-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

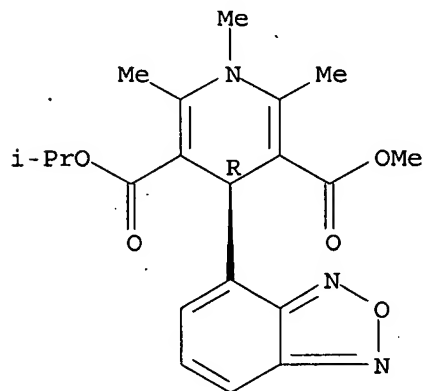
10/564,804



RN 89016-39-7 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-1,2,6-trimethyl-, methyl 1-methylethyl ester, (R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L9 ANSWER 8 OF 8 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1980:639420 CAPLUS

DOCUMENT NUMBER: 93:239420

TITLE: 1,4-Dihydropyridines and their use

INVENTOR(S): Neumann, Peter

PATENT ASSIGNEE(S): Sandoz-Patent-G.m.b.H., Switz.

SOURCE: Ger. Offen., 12 pp.

CODEN: GWXXBX

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT: 4

PATENT INFORMATION:

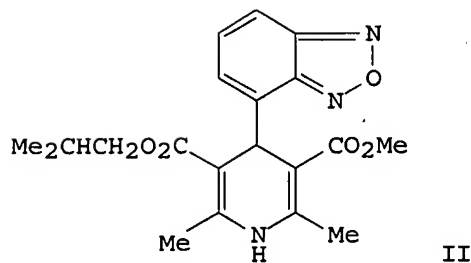
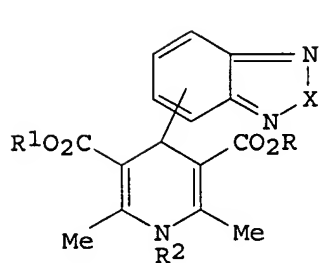
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 2949491	A1	19800626	DE 1979-2949491	19791208
DE 2949491	C2	19881027		
CH 639659	A5	19831130	CH 1978-12835	19781218
SE 7910188	A	19800619	SE 1979-10188	19791211
SE 445219	B	19860609		
SE 445219	C	19860918		
BE 880591	A1	19800613	BE 1979-9643	19791213
NL 7909024	A	19800620	NL 1979-9024	19791214
NL 193066	B	19980506		

NL 193066	C	19980908		
GB 2037766	A	19800716	GB 1979-43113	19791214
GB 2037766	B	19830216		
JP 55083783	A	19800624	JP 1979-163965	19791217
JP 03069910	B	19911105		
AU 7953896	A	19800626	AU 1979-53896	19791217
AU 536055	B2	19840419		
FR 2444681	A1	19800718	FR 1979-30829	19791217
FR 2444681	B1	19821029		
ZA 7906842	A	19810729	ZA 1979-6842	19791218
CH 654836	A5	19860314	CH 1980-5484	19800717
AU 536069	B2	19840419	AU 1982-79464	19820112
AU 8279464	A	19820408		
US 4466972	A	19840821	US 1982-359751	19820319
GB 2103203	A	19830216	GB 1982-15988	19820601
GB 2103203	B	19830608		

PRIORITY APPLN. INFO.:

CH 1978-12835	A	19781218
CH 1977-7520	A	19770620
CH 1978-2865	A	19780316
US 1978-915858	A2	19780615
CH 1978-12888	A	19781218
CH 1978-12890	A	19781218
CH 1979-3472	A	19790411
CH 1979-3477	A	19790411
CH 1979-5627	A	19790615
GB 1979-40624		19791123
DE 1979-2949491	A	19791208
US 1979-101591	A2	19791210
GB 1979-43113	A3	19791214
US 1980-173305	A1	19800729

GI



AB Dihydropyridines I (R = C1-4 alkyl, alkoxyalkyl, cyclopentyl; R1 = C1-4 alkyl; R2 = H, Me, Pr; X = O, S) were prepared for use as vasodilators, antihypertensives, Ca antagonists, and in the treatment of angina pectoris (no data). Thus 2,1,3-benzoxadiazole-4-carboxaldehyde was condensed with AcCH2CO2CH2CHMe2 and H2NCMe:CHCO2Me to give II.

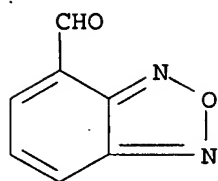
IT 32863-32-4

RL: RCT (Reactant); RACT (Reactant or reagent)
(cyclocondensation of, with acetoacetate and aminocrotonate)

RN 32863-32-4 CAPLUS

CN 2,1,3-Benzoxadiazole-4-carboxaldehyde (CA INDEX NAME)

10/564,804

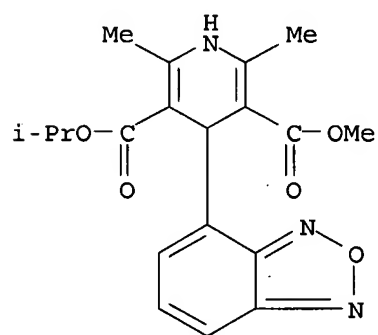


IT 75695-93-1P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

RN 75695-93-1 CAPLUS

CN 3,5-Pyridinedicarboxylic acid, 4-(2,1,3-benzoxadiazol-4-yl)-1,4-dihydro-
2,6-dimethyl-, 3-methyl 5-(1-methylethyl) ester (CA INDEX NAME)



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